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110TH CONGRESS  
2D SESSION

# S. 3164

To amend title XVIII of the Social Security Act to reduce fraud under the Medicare program.

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IN THE SENATE OF THE UNITED STATES

JUNE 19, 2008

Mr. MARTINEZ (for himself and Mr. CORNYN) introduced the following bill; which was read twice and referred to the Committee on Finance

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## A BILL

To amend title XVIII of the Social Security Act to reduce fraud under the Medicare program.

1       *Be it enacted by the Senate and House of Representatives of the United States of America in Congress assembled,*

3 **SECTION 1. SHORT TITLE.**

4       This Act may be cited as the “Seniors and Taxpayers  
5       Obligation Protection Act of 2008”.

1 SEC. 2. REQUIRING THE SECRETARY OF HEALTH AND  
2 HUMAN SERVICES TO CHANGE THE MEDI-  
3 CARE BENEFICIARY IDENTIFIER USED TO  
4 IDENTIFY MEDICARE BENEFICIARIES UNDER  
5 THE MEDICARE PROGRAM.

6 (a) PROCEDURES.—

7 (1) IN GENERAL.—Not later than 1 year after  
8 the date of enactment of this Act, the Secretary of  
9 Health and Human Services (in this section referred  
10 to as the “Secretary”) shall establish and implement  
11 procedures to change the Medicare beneficiary iden-  
12 tifier used to identify individuals entitled to benefits  
13 under part A of title XVIII of the Social Security  
14 Act or enrolled under part B of such title so that  
15 such an individual’s social security account number  
16 is not used.

17 (2) MAINTAINING EXISTING HICN STRUC-  
18 TURE.—In order to minimize the impact of the  
19 change under paragraph (1) on systems that com-  
20 municate with Medicare beneficiary eligibility sys-  
21 tems, the procedures under paragraph (1) shall pro-  
22 vide that the new Medicare beneficiary identifier  
23 maintain the existing Health Insurance Claim Num-  
24 ber structure.

25 (3) PROTECTION AGAINST FRAUD.—The proce-  
26 dures under paragraph (1) shall provide for a proc-

1       ess for changing the Medicare beneficiary identifier  
2       for an individual to a different identifier in the case  
3       of the discovery of fraud, including identity theft.

4                     (4) PHASE-IN AUTHORITY.—

5                         (A) IN GENERAL.—Subject to subparagraphs (B) and (C), the Secretary may phase in  
6       the change under paragraph (1) in such manner as the Secretary determines appropriate.

7                         (B) LIMIT.—The phase-in period under  
8       subparagraph (A) shall not exceed 10 years.

9                         (C) NEWLY ENTITLED AND ENROLLED INDIVIDUALS.—The Secretary shall ensure that  
10      the change under paragraph (1) is implemented not later than January 1, 2010 with respect to  
11      any individual who first becomes entitled to  
12      benefits under part A of title XVIII of the Social Security Act or enrolled under part B of  
13      such title on or after such date.

14                         (b) EDUCATION AND OUTREACH.—The Secretary  
15      shall establish a program of education and outreach for  
16      individuals entitled to benefits under part A of title XVIII  
17      of the Social Security Act or enrolled under part B of such  
18      title, providers of services (as defined in subsection (u) of  
19      section 1861 of such Act (42 U.S.C. 1395x)), and sup-

1 pliers (as defined in subsection (d) of such section) on the  
2 change under paragraph (1).

3 (c) AUTHORIZATION OF APPROPRIATIONS.—There  
4 are authorized to be appropriated such sums as may be  
5 necessary to carry out this section.

6 SEC. 3. MONTHLY VERIFICATION OF ACCURACY OF  
7 CHARGES FOR PHYSICIANS' SERVICES.

8 (a) IN GENERAL.—Section 1893 of the Social Secu-  
9 rity Act (42 U.S.C. 1395ddd) is amended—

10 (1) in subsection (b), by adding at the end the  
11 following new paragraph:

12 “(7) The monthly verification of the accuracy of  
13 charges for physicians' services under the system  
14 under subsection (i).”;

15 (2) in subsection (e), by adding at the end of  
16 the flush matter following paragraph (4), the fol-  
17 lowing new sentence: “In the case of the activity de-  
18 scribed in subsection (b)(7), an entity shall only be  
19 eligible to enter into a contract under the Program  
20 to carry out the activity if the entity is a medicare  
21 administrative contractor with a contract under sec-  
22 tion 1874A.”; and

23 (3) by adding at the end the following new sub-  
24 section:

1       “(i) MONTHLY VERIFICATION OF ACCURACY OF  
2 CHARGES FOR PHYSICIANS’ SERVICES.—

3           “(1) SYSTEM.—

4              “(A) IN GENERAL.—Not later than 1 year  
5 after the date of the enactment of this sub-  
6 section, the Secretary shall establish and imple-  
7 ment a system to verify (electronically or other-  
8 wise, taking into consideration the administra-  
9 tive burden of such verification on physicians  
10 and group practices) on a monthly basis that  
11 the claims for reimbursement under part B for  
12 physicians’ services furnished in high risk areas  
13 are—

14              “(i) for physicians’ services actually  
15 furnished by the physician (or the physi-  
16 cian’s group practice); and

17              “(ii) otherwise accurate.

18            “(B) NO DETERMINATION OF MEDICAL  
19 NECESSITY.—In no case shall any verification  
20 conducted under the system established under  
21 subparagraph (A) include a determination of  
22 the medical necessity of the physicians’ service.

23            “(2) VERIFICATION.—Under the system, the  
24 Secretary, at the end of each month, shall provide  
25 the physician (or the group practice) with a detailed

1       list of such claims for reimbursement that were sub-  
2       mitted during the month in order for the physician  
3       (or the group practice) to review and verify the list.  
4       In providing the detailed list, the Secretary shall use  
5       the provider number of the physician (or the group  
6       practice).

7           “(3) AUDITS.—The Secretary shall conduct au-  
8       dits of the review and verification by physicians and  
9       group practices of the detailed list provided under  
10      paragraph (2). Such audits shall assess whether the  
11      physician or group practice conducted such review  
12      and verification in a fraudulent manner. In the case  
13      where the Secretary determines such review and  
14      verification was conducted in a fraudulent manner,  
15      the Secretary shall recoup any payments resulting  
16      from the fraudulent review and verification and im-  
17      pose a civil money penalty in an amount determined  
18      appropriate by the Secretary on the physician or  
19      group practice who conducted the fraudulent review  
20      and verification. The provisions of section 1128A  
21      (other than subsections (a) and (b)) shall apply to  
22      a civil money penalty under the previous sentence in  
23      the same manner as such provisions apply to a pen-  
24      alty or proceeding under section 1128A(a).

1           “(4) HIGH RISK AREAS DEFINED.—In this sub-  
2       section, the term ‘high risk area’ means a county  
3       designated as a high risk area under subsection  
4       (j)(1).

5           “(5) ACTIONS THROUGH MEDICARE ADMINIS-  
6       TRATIVE CONTRACTORS.—In carrying out this sub-  
7       section, the Secretary shall act through medicare ad-  
8       ministrative contractors with a contract under sec-  
9       tion 1874A.

10          “(6) REPORT BY THE SECRETARY.—Not later  
11       than 1 year after implementation of the system es-  
12       tablished under paragraph (1), the Secretary shall  
13       submit a report to Congress on the progress of such  
14       implementation. Such report shall include rec-  
15       ommendations—

16           “(A) on how to improve such implemen-  
17       tation, including whether the system should be ex-  
18       panded to include verification of claims for re-  
19       imbursement under part B for physicians’ serv-  
20       ices furnished in additional areas; and

21           “(B) for such legislation and administra-  
22       tive action as the Secretary determines appro-  
23       priate.”.

24          (b) AUTHORIZATION OF APPROPRIATIONS.—To carry  
25       out the amendments made by this section, there are au-

1 thorized to be appropriated such sums as may be nec-  
2 essary for each of fiscal years 2009 through 2013.

3 **SEC. 4. DETECTION OF MEDICARE FRAUD IN HIGH RISK**

4 **AREAS.**

5 (a) IN GENERAL.—Section 1893 of the Social Secu-  
6 rity Act (42 U.S.C. 1395ddd), as amended by section 3,  
7 is amended—

8 (1) in subsection (b), by adding at the end the  
9 following new paragraph:

10 “(8) Implementation of prepayment fraud de-  
11 tection methods under subsection (j).”;

12 (2) in subsection (c), in the second sentence of  
13 the flush matter following paragraph (4), by striking  
14 “activity described in subsection (b)(7)” and insert-  
15 ing “activities described in paragraphs (7) and (8)  
16 of subsection (b)”; and

17 (3) by adding at the end the following new sub-  
18 section:

19 “(j) DETECTION OF MEDICARE FRAUD IN HIGH  
20 RISK AREAS.—

21 “(1) ESTABLISHMENT OF SYSTEM TO IDENTIFY  
22 COUNTIES MOST VULNERABLE TO FRAUD.—Not  
23 later than 6 months after the date of the enactment  
24 of this subsection, the Secretary shall establish a  
25 system to identify the 50 counties most vulnerable to

1       fraud with respect to items and services furnished by  
2       providers of services (other than hospitals and crit-  
3       ical access hospitals) and suppliers based on the de-  
4       gree of county-specific reimbursement and analysis  
5       of payment trends under this title. The Secretary  
6       shall designate the counties identified under the pre-  
7       ceding sentence as 'high risk areas'.

8                 “(2) PREPAYMENT FRAUD DETECTION.—The  
9       Secretary shall establish procedures for the imple-  
10      mentation of prepayment fraud detection methods  
11      under this title with respect to items and services  
12      furnished by such providers of services and suppliers  
13      in high risk areas designated under paragraph (1),  
14      including the following:

15                 “(A) Pre-enrollment site visits for such  
16      providers of services and suppliers which have  
17      the highest probability of committing fraud  
18      under this title.

19                 “(B) Data analysis to establish prepay-  
20      ment claim edits designed to target the claims  
21      for reimbursement under this title for such  
22      items and services that are most likely to be  
23      fraudulent.

24                 “(C) Prepayment benefit integrity reviews  
25      for claims for reimbursement under this title

for such items and services that are suspended  
as a result of such edits.

3                 “(3) ACTIONS THROUGH MEDICARE ADMINIS-  
4                 TRATIVE CONTRACTORS.—In carrying out this sub-  
5                 section, the Secretary shall act through medicare ad-  
6                 ministrative contractors with a contract under sec-  
7                 tion 1874A.

8                 “(4) REPORT TO CONGRESS.—The Secretary  
9         shall, upon request, appear and testify before Con-  
10         gress regarding the status of the implementation of  
11         prepayment fraud detection methods under this sub-  
12         section.”.

13       (b) AUTHORIZATION OF APPROPRIATIONS.—To carry  
14 out the amendments made by this section, there are au-  
15 thorized to be appropriated such sums as may be nec-  
16 essary, not to exceed \$50,000,000, for each of fiscal years  
17 2009 through 2013.

18 SEC. 5. STUDY ON THE USE OF TECHNOLOGY FOR REAL-  
19 TIME DATA REVIEW.

(a) STUDY ON THE USE OF TECHNOLOGY FOR REAL-TIME DATA REVIEW.—The Secretary of Health and Human Services shall conduct a study on the use of technology (similar to that used with respect to the analysis of credit card charging patterns) to provide real-time data analysis of claims for reimbursement under the Medicare

1 program under title XVIII of the Social Security Act to  
2 identify and investigate unusual billing or order practices  
3 under the Medicare program that could indicate fraud or  
4 abuse. Such study shall include an analysis of the fol-  
5 lowing:

6               (1) Whether such technology could be used to  
7 identify unusual billing or order practices under the  
8 Medicare program by an individual provider of serv-  
9 ices or for a certain HCPCS code in a particular  
10 area of the country without alerting potentially  
11 fraudulent providers of services and allowing them to  
12 escape or go unnoticed.

13               (2) How such technology can be implemented  
14 under the Medicare program to provide for the effec-  
15 tive review of claim logs in an accurate and timely  
16 manner.

17               (b) REPORT.—Not later than 1 year after the date  
18 of enactment of this Act, the Secretary shall submit a re-  
19 port to Congress on the study conducted under subsection  
20 (a), together with recommendations for such legislation  
21 and administrative action as the Secretary determines ap-  
22 propriate.

1 SEC. 6. EDITS ON 855S MEDICARE ENROLLMENT APPLICA-  
2 TION.

3 Section 1834(a) of the Social Security Act (42 U.S.C.  
4 1395m(a)) is amended by adding at the end the following  
5 new paragraph:

6 “(22) CONFIRMATION WITH NATIONAL SUP-  
7 PLIER CLEARINGHOUSE PRIOR TO REIMBURSE-  
8 MENT.—

9 “(A) IN GENERAL.—Not later than 1 year  
10 after the date of enactment of this paragraph,  
11 the Secretary shall establish procedures to re-  
12 quire carriers, prior to paying a claim for reim-  
13 bursement for durable medical equipment, pros-  
14 thetics, orthotics, and supplies under this title,  
15 to confirm with the National Supplier Clearing-  
16 house—

17 “(i) that the Medicare identification  
18 number of the supplier is active; and

19 “(ii) that the item or service for which  
20 the claim for reimbursement is submitted  
21 was properly identified on the CMS-855S  
22 Medicare enrollment application.

23 “(B) ONLINE DATABASE FOR IMPLEMEN-  
24 TATION.—Not later than 18 months after the  
25 date of enactment of this paragraph, the Sec-  
26 retary shall establish an online database similar

1 to that used for the National Provider Identifier  
2 to enable providers of services, accreditors, car-  
3 riers, and the National Supplier Clearinghouse  
4 to view information on specialties and the types  
5 of items and services each supplier has indi-  
6 cated on the CMS-855S Medicare enrollment  
7 application submitted by the supplier.

8           “(C) NOTIFICATION OF CLAIM DENIAL  
9 AND RESUBMISSION.—In the case where a claim  
10 for reimbursement for durable medical equip-  
11 ment, prosthetics, orthotics, and supplies under  
12 this title is denied because the item or service  
13 furnished does not correctly match up with the  
14 information on file with the National Supplier  
15 Clearinghouse—

16           “(i) the National Supplier Clearing-  
17 house shall—

18           “(I) provide the supplier written  
19 notification of the reason for such de-  
20 nial; and

21           “(II) allow the supplier 60 days  
22 to provide the National Supplier  
23 Clearinghouse with appropriate certifi-  
24 cation, licensing, or accreditation; and

1                         “(ii) the Secretary shall waive applica-  
2                         ble requirements relating to the time frame  
3                         for the submission of claims for payment  
4                         under this title in order to permit the re-  
5                         submission of such claim if payment of  
6                         such claim would otherwise be allowed  
7                         under this title.”.

8     **SEC. 7. SERIAL NUMBER TRACKING SYSTEM FOR DURABLE**  
9                         **MEDICAL EQUIPMENT.**

10                         (a) IN GENERAL.—Section 1834(a) of the Social Se-  
11                         curity Act (42 U.S.C. 1395m(a)), as amended by section  
12                         6(a), is amended by adding at the end the following new  
13                         paragraph:

14                         “(23) SERIAL NUMBER TRACKING SYSTEM FOR  
15                         DURABLE MEDICAL EQUIPMENT.—

16                         “(A) ESTABLISHMENT.—In the case of  
17                         any item of durable medical equipment which  
18                         has not been issued a unique identifier under  
19                         the unique device identification system estab-  
20                         lished under section 519(f) of the Federal  
21                         Food, Drug, and Cosmetic Act, the Secretary  
22                         shall promulgate regulations establishing a sys-  
23                         tem for such durable medical equipment requir-  
24                         ing the label of such equipment to bear a  
25                         unique identifier, unless the Secretary requires

1           an alternative placement or provides an excep-  
2           tion for a particular item or type of durable  
3           medical equipment under such section 519(f).

4           “(B) PROVISION OF UNIQUE IDENTIFIER  
5           TO THE SECRETARY.—A manufacturer of an  
6           item of durable medical equipment shall submit  
7           to the Secretary the unique identifier issued  
8           under subparagraph (A) or such section 519(f)  
9           with respect to such item (in accordance with  
10          procedures established by the Secretary). The  
11          Secretary shall provide for the storage of such  
12          unique identifier in accordance with subpara-  
13          graph (D)(i).

14           “(C) REQUIREMENTS FOR MANUFAC-  
15          TURERS AND WHOLESALERS.—A manufacturer of  
16          an item of durable medical equipment, or, in  
17          the case where a wholesaler provides an item of  
18          durable medical equipment to a supplier, the  
19          wholesaler, shall—

20           “(i) upon issuing an item to a sup-  
21          plier, develop a product description for the  
22          item which includes—

23           “(I) the unique identifier of the  
24          item;

1                         “(II) the specific Healthcare  
2                         Common Procedure Coding System  
3                         (HCPCS) code for the item;

4                         “(III) the name of the supplier  
5                         the item was shipped to; and

6                         “(IV) the supplier’s Medicare  
7                         identification number; and

8                         “(ii) submit the product description  
9                         developed under clause (i) to the Secretary  
10                         for storage in the unique identifier data-  
11                         base in accordance with subparagraph  
12                         (E)(i).

13                         “(D) REQUIREMENTS FOR SUPPLIERS.—A  
14                         supplier of an item of durable medical equip-  
15                         ment shall—

16                         “(i) upon issuing the item to a bene-  
17                         ficiary, note the unique identifier of such  
18                         item on—

19                         “(I) the claim form submitted for  
20                         such item; and

21                         “(II) when appropriate or other-  
22                         wise required, the detailed product de-  
23                         scription of the item;

24                         “(ii) in the case where the item is  
25                         issued to a beneficiary on a rental basis,

1           designate the unique identifier with an 'R'  
2           after the number to indicate that the item  
3           was rented, and not purchased, by the ben-  
4           eficiary; and

5                 “(iii) upon return of the item to the  
6           supplier, notify the Secretary—

7                     “(I) before reissuing that item  
8           and resubmitting that number on  
9           such a claim form; or

10                  “(II) upon resubmitting that  
11           number on such a claim form.

12                 “(E) REQUIREMENTS FOR THE SEC-  
13           RETARY.—

14                 “(i) MAINTENANCE OF DATABASE OF  
15           SERIAL NUMBERS.—The Secretary shall  
16           establish and maintain a database con-  
17           taining the unique identifiers submitted by  
18           manufacturers of items of durable medical  
19           equipment under subparagraph (B).

20                 “(ii) PAYMENT.—

21                 “(I) LIMITATION.—Subject to  
22           subclause (II), payment may only be  
23           made for an item of durable medical  
24           equipment under this part if the  
25           unique identifier on the claim form

1 submitted for such item matches the  
2 unique identifier submitted by the  
3 manufacturer of such item under sub-  
4 paragraph (B).

5 “(II) EXCEPTION TO LIMITATION  
6 AFTER VERIFICATION OF RECEIPT.—  
7 In the case where the unique identi-  
8 fier is not on the claim form sub-  
9 mitted for such item or does not  
10 match the unique identifier submitted  
11 by the manufacturer of such item  
12 under subparagraph (B), no payment  
13 shall be made under this part for the  
14 item of durable medical equipment  
15 until the Secretary has verified that  
16 the beneficiary has received such item  
17 in accordance with subclause (IV).

18 “(III) DUPLICATIVE UNIQUE  
19 IDENTIFIERS.—In the case where a  
20 unique identifier is submitted on more  
21 than 1 claim form submitted for such  
22 an item and there is no indication  
23 from the supplier that the item of du-  
24 rable medical equipment has been re-  
25 turned by 1 beneficiary and is now

1 being used by another beneficiary, no  
2 payment shall be made under this  
3 part for such item of durable medical  
4 equipment unless the Secretary has  
5 verified that the beneficiary has re-  
6 ceived such item in accordance with  
7 subclause (IV).

8 “(IV) VERIFICATION.—The Sec-  
9 retary shall conduct any verification  
10 required under subclause (II) or (III)  
11 within 30 days after receipt by the  
12 Secretary of the relevant claim form.  
13 In the case where such verification is  
14 not completed within such time pe-  
15 riod, the Secretary shall pay such  
16 claim, complete the verification, and,  
17 in the case where the Secretary has  
18 entered into a contract with an entity  
19 for the conduct of such verification,  
20 recover any payments that would not  
21 have been made if the verification had  
22 been completed within such time pe-  
23 riod from such entity.

24 “(iii) QUALITY CONTROL AUDITS.—

25 The Secretary shall conduct quality control

1                   audits to identify unusual billing patterns  
2                   with respect to items of durable medical  
3                   equipment for which payment is made  
4                   under this part and may conduct unan-  
5                   nounced site visits or commission other  
6                   agencies to conduct such site visits as part  
7                   of such quality control audits.

8                   “(iv) NO USE AS A PRECERTIFICATION  
9                   MECHANISM.—In no case shall a unique  
10                  identifier issued under subparagraph (A)  
11                  or section 519(f) of the Federal Food,  
12                  Drug, and Cosmetic Act be used as a  
13                  precertification mechanism for the supply  
14                  of an item of durable medical equipment or  
15                  the payment of a claim for such an item  
16                  under this part.”.

17                 (b) EFFECTIVE DATE.—The amendment made by  
18 subsection (a) shall take effect 3 years after the date of  
19 enactment of this Act.

20 **SEC. 8. SENSE OF THE SENATE REGARDING SURETY BOND**  
21                   **REQUIREMENTS FOR SUPPLIERS OF DURA-**  
22                   **BLE MEDICAL EQUIPMENT.**

23                 (a) FINDINGS.—The Senate finds the following:  
24                   (1) Documented fraud in the Medicare Durable  
25                  Medical Equipment, Prosthetics, Orthotics, and Sup-

1 plies Competitive Bidding Program under section  
2 1847 of the Social Security Act (42 U.S.C. 1395w-  
3) has potentially cost taxpayers in the United  
4 States billions of dollars.

5 (2) Congress, having previously recognized  
6 fraudulent practices with respect to durable medical  
7 equipment under the Medicare program under title  
8 XVIII of the Social Security Act, directed the Sec-  
9 retary of Health and Human Services to take action  
10 against such fraudulent practices through the imple-  
11 mentation of a surety bond requirement under sec-  
12 tion 1834(a)(16) of the Social Security Act (42  
13 U.S.C. 1395m(a)(16)), as added by section 4312 of  
14 the Balanced Budget Act of 1997 (Public Law 105-  
15 33).

16 (3) Such surety bond requirement is necessary  
17 to—

18 (A) limit the risk to the Medicare program  
19 of fraudulent suppliers of durable medical  
20 equipment;

21 (B) enhance the enrollment process under  
22 the Medicare program to ensure that only legiti-  
23 mate suppliers of durable medical equipment  
24 are enrolled or are allowed to remain enrolled in

1                   any programs established or implemented under  
2                   the Medicare program;

3                   (C) ensure that the Medicare program re-  
4                   couples erroneous payments that result from  
5                   fraudulent or abusive billing practices by allow-  
6                   ing the Centers for Medicare & Medicaid Serv-  
7                   ices, or entities under a contract with the Cen-  
8                   ters for Medicare & Medicaid Services, to seek  
9                   payments from a surety up to the penal sum;  
10                  and

11                  (D) help ensure that beneficiaries under  
12                  the Medicare program receive items and serv-  
13                  ices that are considered reasonable and nec-  
14                  essary from legitimate suppliers of durable  
15                  medical equipment.

16                  (4) To date, more than a decade after the en-  
17                  actment of the Balanced Budget Act of 1997 (Public  
18                  Law 105-33), such section 1834(a)(16) has yet to  
19                  be implemented by the Secretary of Health and  
20                  Human Services, potentially costing taxpayers and  
21                  Medicare beneficiaries billions of additional dollars  
22                  and negatively impacting responsible suppliers of du-  
23                  rable medical equipment under the Medicare pro-  
24                  gram.

1       (b) SENSE OF THE SENATE.—It is the Sense of the  
2 Senate that the Secretary of Health and Human Services  
3 must put in place the surety bond requirement under sec-  
4 tion 1834(a)(16) of the Social Security Act (42 U.S.C.  
5 1395m(a)(16)) within 6 months of the date of enactment  
6 of this Act in order to maintain integrity under the Medi-  
7 care program.

○

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